

(19) World Intellectual Property Organization  
International Bureau(43) International Publication Date  
2 August 2001 (02.08.2001)

PCT

(10) International Publication Number  
WO 01/55301 A2

(51) International Patent Classification <sup>7</sup> :	C12N	60/226,681	22 August 2000 (22.08.2000)	US	
		60/227,009	23 August 2000 (23.08.2000)	US	
(21) International Application Number:	PCT/US01/01239	60/228,924	30 August 2000 (30.08.2000)	US	
		60/229,344	1 September 2000 (01.09.2000)	US	
(22) International Filing Date:	17 January 2001 (17.01.2001)	60/229,343	1 September 2000 (01.09.2000)	US	
		60/229,287	1 September 2000 (01.09.2000)	US	
(25) Filing Language:	English	60/229,345	1 September 2000 (01.09.2000)	US	
		60/229,513	5 September 2000 (05.09.2000)	US	
(26) Publication Language:	English	60/229,509	5 September 2000 (05.09.2000)	US	
		60/230,438	6 September 2000 (06.09.2000)	US	
		60/230,437	6 September 2000 (06.09.2000)	US	
(30) Priority Data:		60/231,413	8 September 2000 (08.09.2000)	US	
60/179,065	31 January 2000 (31.01.2000)	US	60/232,080	8 September 2000 (08.09.2000)	US
60/180,628	4 February 2000 (04.02.2000)	US	60/231,414	8 September 2000 (08.09.2000)	US
60/184,664	24 February 2000 (24.02.2000)	US	60/231,244	8 September 2000 (08.09.2000)	US
60/186,350	2 March 2000 (02.03.2000)	US	60/232,081	8 September 2000 (08.09.2000)	US
60/189,874	16 March 2000 (16.03.2000)	US	60/231,242	8 September 2000 (08.09.2000)	US
60/190,076	17 March 2000 (17.03.2000)	US	60/231,243	8 September 2000 (08.09.2000)	US
60/198,123	18 April 2000 (18.04.2000)	US	60/231,968	12 September 2000 (12.09.2000)	US
60/205,515	19 May 2000 (19.05.2000)	US	60/232,401	14 September 2000 (14.09.2000)	US
60/209,467	7 June 2000 (07.06.2000)	US	60/232,399	14 September 2000 (14.09.2000)	US
60/214,886	28 June 2000 (28.06.2000)	US	60/232,400	14 September 2000 (14.09.2000)	US
60/215,135	30 June 2000 (30.06.2000)	US	60/232,397	14 September 2000 (14.09.2000)	US
60/216,647	7 July 2000 (07.07.2000)	US	60/233,063	14 September 2000 (14.09.2000)	US
60/216,880	7 July 2000 (07.07.2000)	US	60/233,064	14 September 2000 (14.09.2000)	US
60/217,487	11 July 2000 (11.07.2000)	US	60/233,065	14 September 2000 (14.09.2000)	US
60/217,496	11 July 2000 (11.07.2000)	US	60/232,398	14 September 2000 (14.09.2000)	US
60/218,290	14 July 2000 (14.07.2000)	US	60/234,223	21 September 2000 (21.09.2000)	US
60/220,963	26 July 2000 (26.07.2000)	US	60/234,274	21 September 2000 (21.09.2000)	US
60/220,964	26 July 2000 (26.07.2000)	US	60/234,997	25 September 2000 (25.09.2000)	US
60/225,757	14 August 2000 (14.08.2000)	US	60/234,998	25 September 2000 (25.09.2000)	US
60/225,270	14 August 2000 (14.08.2000)	US	60/235,484	26 September 2000 (26.09.2000)	US
60/225,447	14 August 2000 (14.08.2000)	US	60/235,834	27 September 2000 (27.09.2000)	US
60/225,267	14 August 2000 (14.08.2000)	US	60/235,836	27 September 2000 (27.09.2000)	US
60/225,758	14 August 2000 (14.08.2000)	US	60/236,369	29 September 2000 (29.09.2000)	US
60/225,268	14 August 2000 (14.08.2000)	US	60/236,327	29 September 2000 (29.09.2000)	US
60/224,518	14 August 2000 (14.08.2000)	US	60/236,370	29 September 2000 (29.09.2000)	US
60/224,519	14 August 2000 (14.08.2000)	US	60/236,368	29 September 2000 (29.09.2000)	US
60/225,759	14 August 2000 (14.08.2000)	US	60/236,367	29 September 2000 (29.09.2000)	US
60/225,213	14 August 2000 (14.08.2000)	US	60/237,039	2 October 2000 (02.10.2000)	US
60/225,266	14 August 2000 (14.08.2000)	US	60/237,038	2 October 2000 (02.10.2000)	US
60/225,214	14 August 2000 (14.08.2000)	US	60/237,040	2 October 2000 (02.10.2000)	US
60/226,279	18 August 2000 (18.08.2000)	US	60/237,037	2 October 2000 (02.10.2000)	US
60/226,868	22 August 2000 (22.08.2000)	US	60/236,802	2 October 2000 (02.10.2000)	US
60/227,182	22 August 2000 (22.08.2000)	US	60/239,937	13 October 2000 (13.10.2000)	US

[Continued on next page]

(54) Title: NUCLEIC ACIDS, PROTEINS, AND ANTIBODIES

(57) Abstract: The present invention relates to novel proteins. More specifically, isolated nucleic acid molecules are provided encoding novel polypeptides. Novel polypeptides and antibodies that bind to these polypeptides are provided. Also provided are vectors, host cells, and recombinant and synthetic methods for producing human polynucleotides and/or polypeptides, and antibodies. The invention further relates to diagnostic and therapeutic methods useful for diagnosing, treating, preventing and/or prognosing disorders related to these novel polypeptides. The invention further relates to screening methods for identifying agonists and antagonists of polynucleotides and polypeptides of the invention. The present invention further relates to methods and/or compositions for inhibiting or enhancing the production and function of the polypeptides of the present invention.

antigen. The affinity of the antibody of interest for a particular antigen and the binding off-rates can be determined from the data by scatchard plot analysis. Competition with a second antibody can also be determined using radioimmunoassays. In this case, the antigen is incubated with antibody of interest conjugated to a labeled compound (e.g.,  $^3\text{H}$  or  $^{125}\text{I}$ ) in the presence of increasing amounts of an unlabeled second antibody.

[275] Antibodies of the invention may be characterized using immunocytochemistry methods on cells (e.g., mammalian cells, such as CHO cells) transfected with a vector enabling the expression of an antigen or with vector alone using techniques commonly known in the art. Antibodies that bind antigen transfected cells, but not vector-only transfected cells, are antigen specific.

#### *Therapeutic Uses*

[276] The present invention is further directed to antibody-based therapies which involve administering antibodies of the invention to an animal, preferably a mammal, and most preferably a human, patient for treating one or more of the disclosed diseases, disorders, or conditions. Therapeutic compounds of the invention include, but are not limited to, antibodies of the invention (including fragments, analogs and derivatives thereof as described herein) and nucleic acids encoding antibodies of the invention (including fragments, analogs and derivatives thereof and anti-idiotypic antibodies as described herein). The antibodies of the invention can be used to treat, inhibit or prevent diseases, disorders or conditions associated with aberrant expression and/or activity of a polypeptide of the invention, including, but not limited to, any one or more of the diseases, disorders, or conditions described herein. The treatment and/or prevention of diseases, disorders, or conditions associated with aberrant expression and/or activity of a polypeptide of the invention includes, but is not limited to, alleviating symptoms associated with those diseases, disorders or conditions. Antibodies of the invention may be provided in pharmaceutically acceptable compositions as known in the art or as described herein.

[277] In a specific and preferred embodiment, the present invention is directed to antibody-based therapies which involve administering antibodies of the invention to an animal, preferably a mammal, and most preferably a human, patient for treating one or more diseases, disorders, or conditions, including but not limited to: neural disorders, immune

stem or progenitor cells, e.g., as obtained from bone marrow, umbilical cord blood, peripheral blood, fetal liver, etc.

[295] In a preferred embodiment, the cell used for gene therapy is autologous to the patient.

[296] In an embodiment in which recombinant cells are used in gene therapy, nucleic acid sequences encoding an antibody are introduced into the cells such that they are expressible by the cells or their progeny, and the recombinant cells are then administered *in vivo* for therapeutic effect. In a specific embodiment, stem or progenitor cells are used. Any stem and/or progenitor cells which can be isolated and maintained *in vitro* can potentially be used in accordance with this embodiment of the present invention (see e.g. PCT Publication WO 94/08598; Stemple and Anderson, *Cell* 71:973-985 (1992); Rheinwald, *Meth. Cell Bio.* 21A:229 (1980); and Pittelkow and Scott, *Mayo Clinic Proc.* 61:771 (1986)).

[297] In a specific embodiment, the nucleic acid to be introduced for purposes of gene therapy comprises an inducible promoter operably linked to the coding region, such that expression of the nucleic acid is controllable by the presence or absence of an appropriate inducer of transcription.

#### *Demonstration of Therapeutic or Prophylactic Activity*

[298] The compounds or pharmaceutical compositions of the invention are preferably tested *in vitro*, and then *in vivo* for the desired therapeutic or prophylactic activity, prior to use in humans. For example, *in vitro* assays to demonstrate the therapeutic or prophylactic utility of a compound or pharmaceutical composition include, the effect of a compound on a cell line or a patient tissue sample. The effect of the compound or composition on the cell line and/or tissue sample can be determined utilizing techniques known to those of skill in the art including, but not limited to, rosette formation assays and cell lysis assays. In accordance with the invention, *in vitro* assays which can be used to determine whether administration of a specific compound is indicated, include *in vitro* cell culture assays in which a patient tissue sample is grown in culture, and exposed to or otherwise administered a compound, and the effect of such compound upon the tissue sample is observed.

#### *Therapeutic/Prophylactic Administration and Composition*